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(54) Color binding mechanism for contact lenses.

(57) Novel colored contact lens and methods for preparation thereof are described. Binding of a color coat to the lenses is accomplished by use of compounds containing the functional groups -CH₂-O-R¹, wherein R¹ is hydrogen or C₁ to C₁₆ alkyl, and said -CH₂-O-R¹ groups are attached to a carbon atom that is part of an aromatic ring, or are attached to a nitrogen or oxygen atom.

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COLOR BINDING MECHANISM FOR CONTACT LENSES

U.S. Patent No. 4,668,240 (Loshak) discloses colored contact lenses produced, in its preferred embodiment, from lens polymers and binding polymers containing the functional groups -OH and -COOH and an additional compound containing at least two groups per molecule of the group -NCO (isocyanate). U.S. Patent Application Serial No. 124,724, filed Nov. 24, 1987, now U.S. Patent No. 4,857,072 discloses a 5 color binding mechanism that is different from that of Loshak, but which also, in its preferred embodiment, requires an isocyanate compound.

Although lenses produced in accordance with the Loshak patent's preferred embodiment are an enormous commercial success, their manufacture requires use of an isocyanate compound for binding the color to the lens. The isocyanate compound has several disadvantages. Its vapors are noxious and toxic if 10 inhaled excessively. Thus it must be used with great caution. The isocyanate compound is very reactive chemically with active hydrogen groups such as -OH in the water molecule. Thus its concentrations can be wastefully reduced, if such molecules are present in the production environment. Its great reactivity also gives rise to pot-life problems, after it is added to the color coating compound. Thus, after the isocyanate compound is added to the color coating ink, the viscosity increases rapidly to a point where the ink must be 15 discarded. The changing viscosity makes control of the coating process difficult and affects the quality of the coating.

The present invention, which does not require use of an isocyanate compound, has several advantages over Loshak's preferred embodiment:

1. Handling noxious and toxic isocyanate compounds is not necessary.
- 20 2. It is possible to print hydrophilic lenses that are fully hydrated, which is not believed practical using ink containing isocyanate compounds.
3. The inks, containing no isocyanate compounds, have better pot-life.
4. Ink viscosity is more stable and easier to control.

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SUMMARY OF THE INVENTION

A first aspect of the invention may be summarized as a method for making a colored contact lens 30 comprising the steps of:

a) providing a contact lens constructed of non-hydrophilic polymers having functional groups selected from at least one of -COOH, -OH, and -NH-R, wherein R is hydrogen or C₁ to C₈ alkyl or hydrophilic polymers;

b) coating at least a portion of a surface of the lens with a color coat comprising at least one pigment, 35 binding polymer having functional groups selected from at least one of -COOH, -OH, and -NH-R, wherein R is hydrogen or C₁ to C₈ alkyl, and an adhesion promotor having at least two functional groups per molecule of the formula -CH₂-O-R¹ wherein R¹ is hydrogen or C₁ to C₁₆ alkyl, and said -CH₂-O-R¹ groups are attached to a carbon atom that is part of an aromatic ring, or attached to a nitrogen or oxygen atom;

c) subjecting the coated lens to conditions which cause the color coat to adhere to the lens.

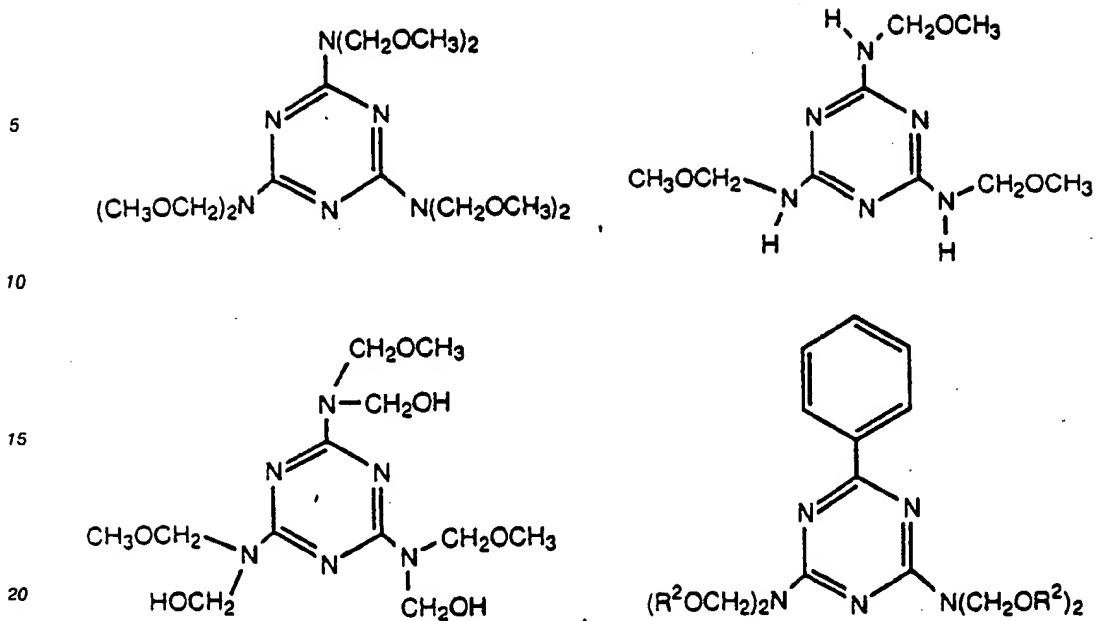
40 Preferably, the lens polymer is hydrophilic and has functional groups selected from at least one of -COOH, -OH, and -NH-R, wherein R is hydrogen or C₁ to C₈ alkyl.

Also preferred is that the lens polymer and binding polymer are hydrophilic and are formed from monomers comprising at least one of acrylic acid, methacrylic acid, hydroxy C₁ to C₆ alkyl ester of acrylic acid and methacrylic acid, amino C₁ to C₈ alkyl ester of acrylic and methacrylic acid, glycerol esters of acrylic and methacrylic acid, and mixtures thereof.

45 More preferably, the lens polymer and binding polymers are formed from a mixture of monomers comprising hydroxyethyl methacrylate, ethoxyethyl methacrylate, and methacrylic acid.

Also preferred is that the lens polymer is hydrophilic and is formed from monomer comprising vinyl pyrrolidone and hydrophobic monomer.

50 Also preferred is that the adhesion promotor is



wherein R² is CH₃ or C₂H₅.

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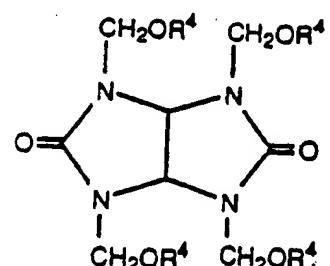
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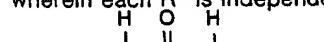
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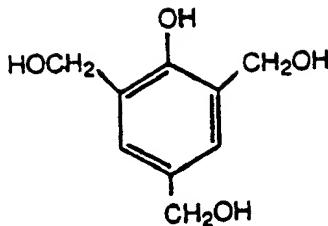
wherein R is a carbon-to-carbon single bond or C₁ to C₆ alkylene and R⁵ is independently C₁ to C₆ alkyl.



wherein each R⁴ is independently H, -CH₃, -C₂H₅, -C₃H₇, or -C₄H₉.



wherein each R⁶ is independently H, -CH₃, or isobutyl, or



10 polymers thereof and combinations thereof.

A second aspect of the invention may be summarized as a method for making a colored contact lens comprising the steps of:

15 a) providing a contact lens constructed of non-hydrophilic polymers having functional groups selected from at least one of -COOH, -OH, and -NH-R, wherein R is hydrogen or C₁ to C₈ alkyl or hydrophilic polymers;

b) coating at least a portion of a surface of the lens with a color coat comprising at least one pigment and adhesion promotor having functional groups of the formula -CH₂-O-R¹ wherein R¹ is hydrogen or C₁ to C₁₆ alkyl with said -CH₂-O-R¹ functional groups attached to a carbon atom that is part of an aromatic ring, or attached to a nitrogen or oxygen atom;

20 c) subjecting the coated lens to conditions which cause the color coat to adhere to the lens.

A third aspect of the invention may be summarized as a method for making a colored contact lens comprising the steps of:

25 a) providing a contact lens constructed of polymer having functional groups of the formula -CH₂-O-R¹ wherein R¹ is hydrogen or C₁ to C₁₆ alkyl with said -CH₂-O-R¹ functional groups attached to a carbon atom that is part of an aromatic ring or attached to a nitrogen or oxygen atom;

b) coating at least a portion of a surface of the lens with a color coat comprising at least one pigment and binding polymer having functional groups selected from at least one of the -COOH, -OH, and -NH-R, wherein R is hydrogen or C₁ to C₈ alkyl, and

c) subjecting the coated lens to conditions which cause the color coat to adhere to the lens.

30 A fourth aspect of the invention may be summarized as a method for making a colored contact lens comprising the steps of:

35 a) providing a contact lens constructed of polymer having functional groups of the formula -CH₂-O-R¹ wherein R¹ is hydrogen or C₁ to C₁₆ alkyl with said -CH₂-O-R¹ functional groups attached to a carbon atom that is part of an aromatic ring or attached to a nitrogen or oxygen atom;

b) coating at least a portion of a surface of the lens with a color coat comprising at least one pigment, binding polymer having functional groups of the formula -CH₂-O-R¹ wherein R¹ is hydrogen or C₁ to C₁₆ alkyl with said -CH₂-O-R¹ functional groups attached to a carbon atom that is part of an aromatic ring or attached to a nitrogen or oxygen atom, and an adhesion promotor having at least two groups per molecule selected from at least one of -COOH, -OH, and -NH-R, wherein R is hydrogen or C₁ to C₈ alkyl, and

40 c) subjecting the coated lens to conditions which cause the color coat to adhere to the lens.

A fifth aspect of the invention may be summarized as a colored contact lens made by any of the four methods described above.

A sixth aspect of the invention may be summarized as a colored contact lens comprising:

45 a) a lens constructed of polymer;

b) pigment mixed with adhesion promotor on at least a portion of a surface of the lens; wherein the adhesion promotor and lens polymer are bonded to each other by the reaction of first functional groups selected from at least one of -COOH, -OH, and -NH-R, wherein R is hydrogen or C₁ to C₈ alkyl with second functional groups of the formula -CH₂-O-R¹ wherein R¹ is hydrogen or C₁ to C₁₆ alkyl with said second functional groups attached to a carbon atom that is part of an aromatic ring or attached to a nitrogen or oxygen atom. Preferably, the lens polymer is formed from at least one monomer selected from acrylic acid, methacrylic acid, hydroxy C₁ to C₈ alkyl ester of acrylic acid or methacrylic acid, amino C₁ to C₈ alkyl ester of acrylic or methacrylic acid, glycerol ester of acrylic or methacrylic acid and mixtures thereof. Most preferably, the lens polymer is formed from at least one monomer selected from acrylic acid, methacrylic acid, and hydroxyethyl methacrylate.

55 A major advantage of this invention is that the compounds used as adhesion promoters or for the dual purpose of pigment binding and adhesion promotion, depend on a condensation, acid-base catalysis mechanism which differs from that of the isocyanate compounds of the prior art. Thus, while the isocyanate compounds are very reactive with even trace amounts of water or alcohol, the compounds of this invention

are generally stable when water or alcohol are present in large quantities. This type of chemical reaction permits greater control of reaction speed and a wider range of compatibility, depending on the specific structure and reaction media, such as water, alcohol, and other active hydrogen compounds which present problems with the isocyanate compounds. The result is as much greater control of pot life and viscosity

5 variation which can translate into improved product quality.

DETAILED DESCRIPTION OF THE INVENTION

10 The lenses colored by the first aspect of this invention may be constructed of non-hydrophilic polymer formed from hydrophobic monomers such as methyl methacrylate, t-butyl styrene (U.S. Patent No. 4,228,269, Loshaek and Shen) or silicone polymer (e.g. U.S. Patent No. 4,120,570, Gaylord) and hydrophilic monomers having functional groups selected from at least one of -COOH, -OH, and -NH-R, wherein R is

15 hydrogen or C₁ to C₈ alkyl. The amount of hydrophilic monomers is not sufficient to render the lens hydrophilic, i.e., the lenses do not absorb a substantial amount of water and form hydrogels.

20 Alternatively the lenses colored by the first aspect of the invention may be constructed of hydrophilic polymer which has the functional groups -COOH, -OH, and/or -NH-R; or the hydrophilic polymer may be devoid of these functional groups. In either case, the hydrophilic polymers absorb substantial amounts of water to form hydrogels. Preferred hydrophilic polymers containing the functional groups are formed from, e.g. acrylic acid, methacrylic acid, hydroxy C₁ to C₆ alkyl esters of acrylic and methacrylic acid, (e.g. hydroxyethyl methacrylate and hydroxybutyl acrylate), amino C₁ to C₈ alkyl esters of acrylic and methacrylic acid (e.g. aminoethyl acrylate or N-ethyl amino methyl methacrylate), glycerol esters of acrylic and methacrylic acid, and mixtures thereof. Preferred hydrophilic lenses having the functional groups are

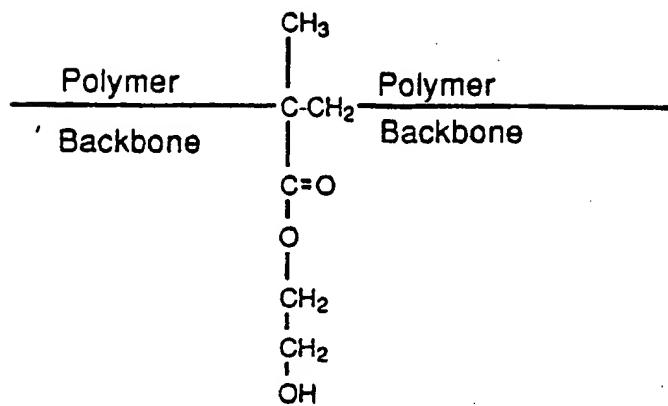
25 disclosed in U.S. Patent No. 4,405,773, (Loshaek) and are formed from a mixture of hydroxyethyl methacrylate, ethoxyethyl methacrylate and methacrylic acid. Preferred hydrophilic monomers not having the functional groups are disclosed in U.S. Patent No. 4,158,089 (Loshaek) and are formed from vinyl pyrrolidone, which is a hydrophilic monomer, and hydrophobic monomers, such as alkyl esters of acrylic and methacrylic acid. Other contact lenses made of hydrophilic polymer that are substantially devoid of the

30 functional groups -COOH, -OH, -NH-R, wherein R is hydrogen or C₁ to C₈ alkyl, -NCO, and epoxy are described in U.S. Patent Application Serial No. 124,724 filed November 24, 1987, now U.S. Patent No. 4,857,072. Typical polymers of this type are produced by copolymerizing a hydrophilic vinyl heterocyclic monomer, such as N-vinyl pyrrolidone, Nsuccinimide, N-vinyl-e-caprolactam, vinyl pyridine, or N-vinyl glutarimide and a hydrophobic monomer such as styrene or C₁ to C₈ alkyl esters of acrylic or methacrylic acid, e.g. methyl methacrylate, ethyl methacrylate, methyl acrylate, etc. The lens polymer is typically cross-linked with a cross-linking agent such as diallyl itaconate, allyl methacrylate, or ethylene glycol dimethacrylate. Typical hydrophilic lenses devoid of the groups -OH, -COOH, and -NH-R, -NCO and epoxy are described in U.S. Patent Nos. 4,158,089 (Loshaek), 4,182,802 (Loshaek), and British patent No. 1,395,501 (National Research Development Corp.).

40 Hydrophilic lenses substantially devoid of the foregoing functional groups colored by this invention are preferably formed from monomers comprising hydrophilic monomer, such as N-vinyl pyrrolidone. The amount of N-vinyl pyrrolidone in the lens is preferably from about 50 to about 90 weight percent, more preferably from about 60 to about 80 weight percent and most preferably from about 65 to about 75 weight percent. If an N-vinyl heterocyclic hydrophilic monomer other than N-vinyl pyrrolidone is used, the above

45 preferred weight percentages also apply. Generally, the amount of hydrophobic monomer employed in a lens is not sufficient to render the lens hydrophobic.

50 The term polymer means a material formed by polymerizing one or more monomeric compounds or a blend of such monomers. For polymers having functional groups, the functional groups project from the polymer backbone or from another group that projects from the backbone. For example, polymerized hydroxyethyl methacrylate may be represented by:



The hydroxyl functional group is at the bottom of the above representation.

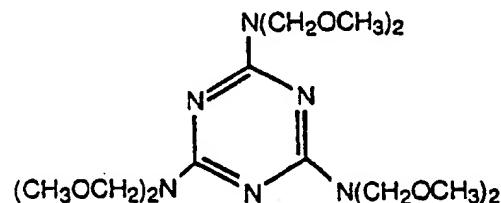
Polymerization and lens shaping are well known in the art and are not part of this invention. Any of the well known techniques may be used.

20 Well known techniques may be used.
25 To produce lenses in accordance with the first aspect of the invention a lens constructed of non-hydrophilic polymers having functional groups selected from at least one of -COOH, -OH, and -NH-R, wherein R is hydrogen or C₁ to C₈ alkyl or hydrophilic polymer is coated on at least a portion of its surface with a color coat comprising at least one pigment, binding polymer having functional groups selected from at least one of -OH, -COOH, and -NH-R, and an adhesion promotor having at least two functional groups of the formula -CH₂-O-R' wherein R' is hydrogen or C₁ to C₁₆ alkyl with said two functional groups per molecule of adhesion promotor attached to a carbon atom that is part of an aromatic ring or attached to a nitrogen or oxygen atom. The description of the binding polymer having the required functional groups is the same as that given previously for lens polymers having the functional groups.
The adhesion promotor having at least two groups of the formula -CH₂-O-R' per molecule of the

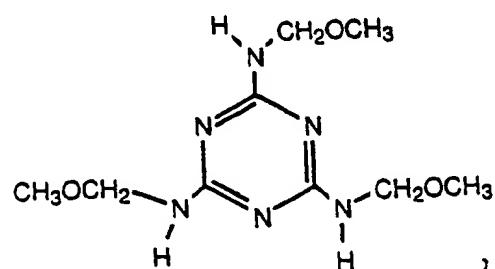
30 The adhesion promotor having at least two groups of the formula $-\text{CH}_2\text{-O-}\text{R}^1$ per molecule of the adhesion promotor are generally formed by reaction of formaldehyde with various other compounds in the ratio of at least two molecules of formaldehyde per molecule of the other compounds. Preferred adhesion promotors, all of which are commercially available are:

A hexamethoxymethylmelamine resin sold as Cymel 301™ having the simplified structural formula

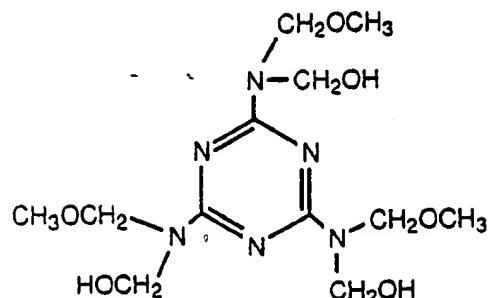
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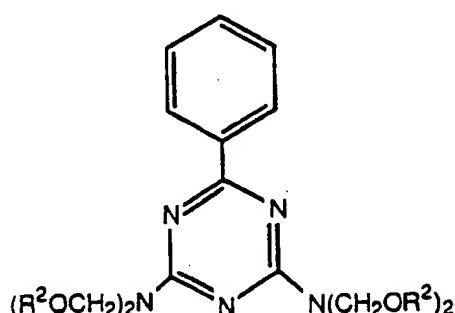
45 A methoxymethyl-imino melamine-formaldehyde resin sold as Cymel 327™ having the simplified structural formula



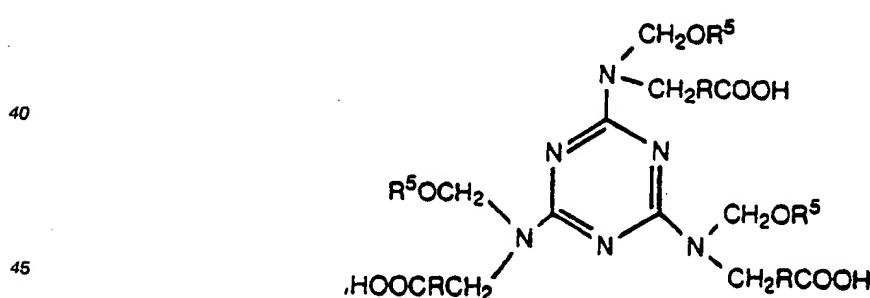
5 A methoxymethyl-methylol melamine-formaldehyde resin sold as Cymel 385TM, having the simplified
10 structural formula



20 A methylated ethylated benzoquanamine having the simplified structural formula



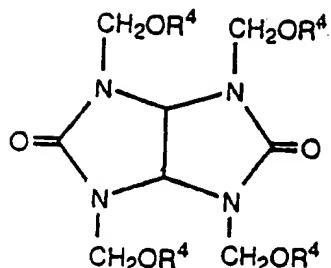
35 wherein each R^2 independently is -CH₃, -CH₂CH₃ or combinations thereof, sold as Cymel 1123TM;
A methylated-isobutylated melamine-formaldehyde resin which contains acidic groups having the simplified
structural formula



50 wherein R is independently a carbon-to-carbon single bond or C₁ to C₆ alkylene preferably methylene, and
R^5 is independently C₁ to C₆, sold as Cymel 1141TM;
A methylated ethylated or tetramethylol glycoluril resin having the simplified structural formula

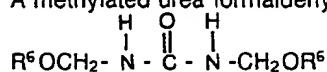
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wherein each R⁴ is independently H, -CH₃, or -C₂H₅ sold as Cymel 1171TM and Cymel 1172TM
 A methylated urea formaldehyde resin having the simplified formula

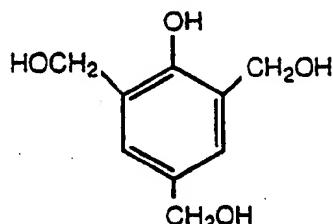
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wherein R⁶ is independently H, -CH₃, or isobutyl sold as Beetle 60TM; and
 Resole resin having the simplified formula

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CymelTM resins and Beetle 60TM resin are available from The American Cyanamid Co., Wayne, New Jersey. Resole resins are available from Monomer-Polymer and Dajac Laboratories, Inc., Trevose, Pennsylvania.

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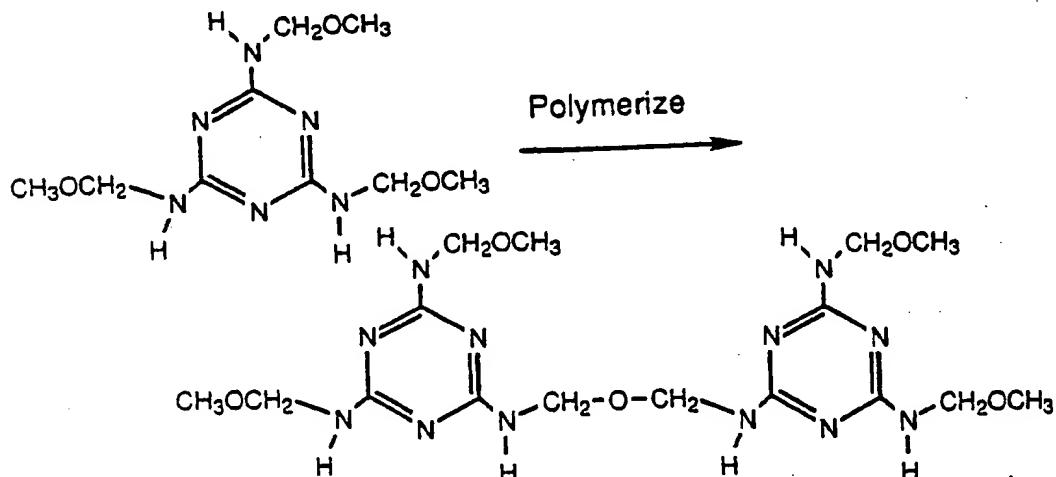
All of the above compounds having at least two groups per molecule of -CH₂-O-R¹ are believed to be polymerized to a minor extent, hence each structural formula is described as being "simplified". For example Cymel 327TM may polymerize according to the reaction:

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Even polymerized, such polymers will still contain at least two groups per molecule of -CH₂-O-R¹.

To achieve reactivity, the methylol (i.e. CH₂OH) or substituted methylol group is attached to an atom of nitrogen, oxygen, or to a carbon atom that is part of an aromatic ring. The aromatic ring is preferably a

benzene ring, as in the resol resin compound, but other unsaturated rings may be used such as, rings of furan, pyrrole, imidazole, thiophene, thiazol, naphthalene, indole, purine, anthracene, phenanthrene, and the like.

To perform the coating step, the mixture of binding polymer having the required functional groups, 5 pigment, additional compound having at least two groups per molecule of $-\text{CH}_2\text{O-R}^1$, and solvent is coated onto at least a portion of the front or rear surface of the lens. Although coating in a single step with this mixture is preferred, the coating step may be performed in two or more substeps, e.g. first coating the lens with a mixture of binding polymer, pigment, and solvent, and thereafter applying a coat of solvent and additional compound over the first coat.

10 The binding polymer, if used or when present, is preferably prepared by the same procedure described in Loshaek's U.S. Patent 4,668,240, column 5, i.e., using a solution that contains the monomers that will be used to form the binding polymers, solvent for the monomer, a molecular weight modifier, and a polymerization initiator. Polymerization preferably is carried out in a heated reaction vessel. It should be noted, however that the solvent used in making the binding polymer, if any, in the present invention need 15 not be constrained to molecules without active hydrogen groups, because of the aforementioned different reaction mechanisms for the adhesion promoter.

20 Suitable molecular weight modifiers include 2-mercaptoethanol, 1-dodecyl mercaptan, other alkyl mercaptans or extra solvent. Suitable polymerization initiators include free radical initiator such as 2,2-azobis(isobutyronitrile), benzoyl peroxide, t-butyl peroxybenzoate, and t-butyl peroxide. Suitable solvents 25 include ethyl cellosolve, cyclopentanone, ethanol, t-butanol, acetone and other aliphatic and alicyclic ketones, ethyl lactate, and monomers used to make the binding polymer.

25 The choice of pigment is very flexible. Any coloring substance or combination thereof that provides a desired color may be used. Preferred pigments include (C.I. is the color index no.) for blue, phthalocyanine green (pigment blue 15, C.I. 74160), cobalt blue (pigment blue 36, C.I. 77343); for green phthalocyanine 30 green (Pigment green 7, C.I. 74260) and chromium sesquioxide; for yellow, red, brown, or black, various iron oxides; for violet, carbazole violet. Of course, blends of such coloring substances are used to achieve the desired shade. For an opaque color coat, titanium dioxide or other opaquing agent may be used. Opaquing agents, which are considered to be pigment within the present specification and claims, are blended with the other pigments into the color coat.

30 After coating, the lens is subjected to conditions that cause the color coat to adhere to the lens, thereby entrapping the pigment within the binding polymer in a manner that forms a colored contact lens. The coated lens, for example, may remain at ambient room temperature where most of the solvents are dried and then cured in an oven that may be purged with inert gas, e.g. nitrogen, and/or be under a partial vacuum. Of course, any conditions that bring about adhesion of the color coat may be used.

35 If the lens is constructed of a hydrophilic polymer, it is hydrated after curing by methods commonly used for uncolored lenses. The concentrations of ingredients in the color coat may vary quite widely, depending on the particular binding polymer and additional compound chosen. The examples provide specific formulations that may be used and the following table gives general ranges:

40 WEIGHT PERCENT OF COMPONENTS IN COLOR COATS		
Ingredient	Broad Range	Preferred Range
Pigment	0.5-30	6.5-20
Binding Polymer (including any residual unreacted monomer)	0-50	13-25
Additional Monomer	0-50	15-30
Adhesion Promotor with ≥ 2 $-\text{CH}_2\text{O-R}^1$ groups	0.5-10	0.5-5
Polymerization Initiator ⁽¹⁾	0-1.0	0.05-0.5
Acid Catalyst	0-10	0-3
Solvent ⁽²⁾	Add to 100 percent	Add to 100 percent

50 ⁽¹⁾ Normally used only if the ink contains additional monomers.

⁽²⁾ Includes solvent from solution polymerization of binding polymers, if solution polymerization was performed and solvents used in pigment grinding.

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The following examples are intended to illustrate, but not limit, the invention. In these examples, the color coat is printed onto the lens and hence is termed "ink". Of course, other types of color coats, e.g. paints, may also be suitable.

PREPARATIVE EXAMPLE NO. 1 CLEAR CONTACT LENSES					
Lens Polymer Identification		L1	L2	L3	
Ingredient		Weight Fraction			
5	Hydroxyethyl Methacrylate (HEMA)	.8838	--	--	
	Ethoxyethyl Methacrylate (EOEMA)	.0982	--	--	
	Methacrylic Acid (MAA)	.0170	--	--	
10	N,N-Dimethylacrylamide	--	--	.7068	
	N-Vinyl Pyrrolidone	--	.7056	--	
	Methyl Methacrylate	--	.2643	.2623	
	Isobutyl Acrylate	--	.0249	.0249	
	Diallyl Itaconate	--	.0016	.0015	
15	Azobisisobutyronitrile	.0010	--	--	
	2,5-Dimethyl-2,5-Bis (2-ethyl hexoylperoxy) Hexane	--	.0030	.0030	
	t-Butyl Peroxybenzoate	--	.0006	.0015	
	<u>TOTAL</u>	<u>1.0000</u>	<u>1.0000</u>	<u>1.0000</u>	

20

Preparation of Clear Lens L1

25 A mixture of all of the ingredients is degassed at about 0.1 millimeters of mercury (mmHg) for about 15 minutes and sealed in a glass tube and the tube is placed in a constant temperature water bath at 35 degrees Centigrade (°C) for about 7 days. The tube is then transferred to an oven and the temperature is raised in steps to 90 °C over a period of about 48 hours. The oven is allowed to cool, and the tube is removed from the oven. The glass tube is broken and a rod of polymer is obtained. The polymer rod is cut into bonnets which are in turn machined into contact lenses by procedures well known in the art.

30

Preparation of Clear Lens L2

35 The ingredients were degassed at about 150 mmHg for about 15 minutes in a flask, then placed in a glass tube. The filled tube was capped and placed in a water bath for 72 hours at 20 °C and then for 96 hours (hrs.) at 30 °C. The ingredients solidified during this time. The glass tube was removed from the water bath and subjected to the following treatment in an oven: 24 hrs. at 50 °C, 4 hrs. at 70 °C and 24 hrs. at 110 °C. The polymer was then removed from the glass tube and heated for 2 hrs. at 140 °C. The polymer was formed into contact lenses by conventional methods.

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Preparation of Clear Lens L3

45 Clear lens L3 is prepared in substantially the same manner as described for clear lens L2 except the filled glass tubes were heated at 35 °C for 6 days in a water bath before subjecting to the oven treatment.

Lens L1, containing polymerized HEMA and MAA, has the functional groups -OH and -COOH. Neither Lens L2 or lens L3 has these functional groups. All three polymers are hydrophilic and swell in saline solution to form a so-called soft lens. However before swelling, they are printed with inks as described later.

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PREPARATIVE EXAMPLE 2 BINDING POLYMER SOLUTION		
5	Solution Identification	S1
	Ingredient	Weight Fraction
10	Hydroxyethyl Methacrylate	.3776
	Ethoxyethyl Methacrylate	.0416
	Methacrylic Acid	.0064
	Cyclopentanone	.5699
	Azobisisobutyronitrile	.0022
	2-Mercaptoethanol	.0022
15	Hydroquinone <u>Monomethylether</u>	.0001
	TOTAL	1.0000

20 Polymerization Procedure

Binder polymerization was conducted by charging all the ingredients except hydroquinone monomethylether to a three neck flask equipped with a stirrer, nitrogen inlet, and an exhaust port. The flask was placed in a water bath held at 40 °C. Nitrogen was passed over the slowly agitated solution.

25 The polymerization was sampled periodically until the viscosity measured between 20,000 and 40,000 centipoise (cps) on a Brookfield Viscometer with a number 3 spindle at 3 revolutions per minute (rpm). Hydroquinone monomethylether was added at this time. The product was then removed from the three neck flask and stored in a refrigerator.

30 In this case the binding polymer was prepared in solution by solution polymerization. Of course other methods of polymerization may be used, e.g. suspension or bulk polymerization.

PREPARATIVE EXAMPLE NO. 3 INKS						
35	Inks Containing Cymel 301TM:					
	Ink Identification	1	2	3	4	
40	Ingredient	Weight Fraction				
	Hydroxyethyl Methacrylate	.2332	.2139	.2234	.2276	
	Ethoxyethyl Methacrylate	.0264	.0242	.0252	.0258	
	Methacrylic Acid	.0043	.0040	.0042	.0042	
	Azobisisobutyronitrile	.0013	.0012	.0013	.0013	
	Polymerized Binder Solution S1	.5034	.4616	.4828	.4956	
	Ethyl Lactate	.1615	.1481	.1549	.1590	
45	Phthalocyanine Blue	.0027	.0025	.0026	.0026	
	Titanium Dioxide	.0551	.0506	.0529	.0543	
	Cymel 301TM*	.0121	.0939	.0527	.0254	
	Catalyst 4040TM**	--	--	--	.0042	
50	TOTAL	1.0000	1.0000	1.0000	1.0000	

*Cymel 301TM is a hexamethoxymethylmelanine resin.

**Catalyst 4040TM is a 40% solution of toluene sulfonic acid in isopropanol added to activation solution.

Inks Containing Cymel 327TM:				
Ink Identification				
	5	6	7	8
Ingredient	Weight Fraction			
Hydroxyethyl Methacrylate	.2345	.2145	.2319	.2140
Ethoxyethyl Methacrylate	.0265	.0243	.0262	.0242
Methacrylic Acid	.0044	.0040	.0040	.0037
Azobisisobutyronitrile	.0013	.0012	.0013	.0012
Polymerized Binder Solution S1	.5037	.4633	.5003	.4616
Ethyl Lactate	.1616	.1487	.1605	.1482
Phthalocyanine Blue	.0027	.0025	.0026	.0024
Titanium Dioxide	.0552	.0508	.0548	.0505
Cymel 327TM*	.0101	.0907	.0184	.0942
Catalyst 4040TM**	--	--	--	--
TOTAL	1.000	1.000	1.000	1.000

*Cymel 327TM is a 90% solution in isobutanol of a methoxymethyl-imino melamine-formaldehyde resin.

**Catalyst 4040TM is a 40% solution of toluene sulfonic acid in isopropanol.

	9	10	11	12
Ingredient	Weight Fraction			
Hydroxyethyl Methacrylate	.2243	.2108	.2212	.2112
Ethoxyethyl Methacrylate	.0254	.0239	.0250	.0239
Methacrylic Acid	.0042	.0039	.0041	.0039
Azobisisobutyronitrile	.0013	.0012	.0012	.0012
Polymerized Binder Solution S1	.4839	.4555	.4827	.4595
Ethyl Lactate	.1554	.1462	.1549	.1475
Phthalocyanine Blue	.0026	.0024	.0026	.0024
Titanium Dioxide	.0530	.0499	.0529	.0503
Cymel 327TM*	.0384	.0954	.0528	.0976
Catalyst 4040TM**	.0115	.0108	.0026	.0025
TOTAL	1.0000	1.0000	1.0000	1.0000

* Cymel 327TM is a 90% solution in isobutanol of methoxymethyl-imino melamine-formaldehyde resin.

** Catalyst 4040TM is a 40% solution of toluene sulfonic acid in isopropanol. In inks 9 and 10, catalyst 4040 was added to the activated ink paste. In inks 11 and 12 catalyst 4040 was added to the activation solution.

Inks Containing Cymel 385TM:		
Ink Identification	13	14
Ingredient	Weight Fraction	
Hydroxyethyl-Methacrylate	.2315	.2128
Ethoxyethyl-Methacrylate	.0261	.0240
Methacrylic Acid	.0043	.0040
Azobisisobutyronitrile	.0013	.0012
Polymerized Binder Solution S1	.5030	.4629
Ethyl Lactate	.1614	.1486
Phthalocyanine Blue	.0027	.0025
Titanium Dioxide	.0551	.0507
Cymel 385TM*	.0119	.0908
Catalyst 4040TM**	.0027	.0025
TOTAL	1.0000	1.0000

* Cymel 385TM is an 80% solution in water of methoxymethyl-methyol melamine-formaldehyde resin.

** Catalyst 4040TM is a 40% solution of toluene sulfonic acid in isopropanol added to the activation solution.

Inks Containing Cymel 1123TM:		
Ink Identification	15	16
Ingredient	Weight Fraction	
Hydroxyethyl Methacrylate	.2311	.2106
Ethoxyethyl Methacrylate	.0261	.0238
Methacrylic Acid	.0043	.0039
Azobisisobutyronitrile	.0013	.0012
Polymerized Binder Solution S1	.4981	.4552
Ethyl Lactate	.1598	.1461
Phthalocyanine Blue	.0027	.0024
Titanium Dioxide	.0546	.0499
Cymel 1123TM*	.0111	.0968
Catalyst 4040TM**	.0109	.0101
TOTAL	1.0000	1.0000

* Cymel 1123TM is a methylated, ethylated benzoguanamine resin.

** Catalyst 4040TM is a 40% solution of toluene sulfonic acid in isopropanol added to the activated ink paste.

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Inks Containing Cymel 1141TM:		
Ink Identification	17	18
Ingredient	Weight Fraction	
Hydroxyethyl Methacrylate	.2307	.2106
Ethoxyethyl Methacrylate	.0261	.0238
Methacrylic Acid	.0043	.0039
Azobisisobutyronitrile	.0013	.0012
Polymerized Binder Solution S1	.4985	.4563
Ethyl Lactate	.1600	.1463
Phthalocyanine Blue	.0026	.0024
Titanium Dioxide	.0546	.0500
Cymel 1141TM*	.0120	.0922
Catalyst 4040TM**	.0099	.0133
<u>TOTAL</u>	1.0000	1.0000

* Cymel 1141TM is an 85% solution in isobutanol of a methylated-isobutylated melamine-formaldehyde resin which contains acidic groups.

** Catalyst 4040TM is a 40% solution of toluene sulfonic acid in isopropanol added to the activated ink paste.

Inks Containing Cymel 1171TM:		
Ink Identification	19	
Ingredient	Weight Fraction	
Hydroxyethyl Methacrylate	.2308	
Ethoxyethyl Methacrylate	.0261	
Methacrylic Acid	.0043	
Azobisisobutyronitrile	.0013	
Polymerized Binder Solution S1	.4963	
Ethyl Lactate	.1593	
Phthalocyanine Blue	.0026	
Titanium Dioxide	.0544	
Cymel 1171TM*	.0147	
Catalyst 4040TM**	.0102	
<u>TOTAL</u>	1.0000	

* Cymel 1171TM is a 95% solution in N-butyl alcohol of a methylated, ethylated glycoluril resin.

** Catalyst 4040TM is a 40% solution of toluene sulfonic acid in isopropanol added to the activated ink paste.

Inks Containing Cymel 1172TM:					
Ink Identification		20	21	22	23
Ingredient		Weight Fraction			
	Hydroxyethyl Methacrylate	.2211	.2062	.2328	.2246
	Ethoxyethyl Methacrylate	.0250	.0233	.0264	.0254
	Methacrylic Acid	.0042	.0039	.0043	.0042
	Azobisisobutyronitrile	.0013	.0012	.0013	.0013
	Polymerized Binder Solution S1	.4768	.4457	.5039	.4843
	Ethyl Lactate	.1530	.1430	.1617	.1554
	Phthalocyanine Blue	.0025	.0024	.0027	.0026
	Titanium Dioxide	.0522	.0488	.0552	.0531
	Cymel 1172TM*	.0490	.0977	.0117	.0491
	Catalyst 4040TM**	.0149	.0278	--	--
	<u>TOTAL</u>	1.0000	1.0000	1.0000	1.0000

* Cymel 1172TM is a 45% solution in water of tetramethylol glycoluril.

** Catalyst 4040TM is a 40% solution of toluene sulfonic acid in isopropanol added to the activated ink paste.

Inks Containing Beetle 60TM:					
Ink Identification		25	26	27	28
Ingredient		Weight Fraction			
	Hydroxyethyl Methacrylate	.2305	.2240	.2143	.2285
	Ethoxyethyl Methacrylate	.0261	.0254	.0243	.0258
	Methacrylic Acid	.0043	.0042	.0040	.0042
	Azobisisobutyronitrile	.0013	.0013	.0012	.0013
	Polymerized Binder Solution S1	.4975	.4839	.4635	.4941
	Ethyl Lactate	.1597	.1552	.1487	.1585
	Phthalocyanine Blue	.0026	.0026	.0025	.0026
	Titanium Dioxide	.0545	.0530	.0508	.0541
	Beetle 60TM*	.0235	.0504	.0907	.0309
	Catalyst 4040TM**	--	--	--	.0042
	<u>TOTAL</u>	1.0000	1.0000	1.0000	1.0000

* Beetle 60TM is a 85% solution in isopropyl alcohol of a methylated urea-formaldehyde resin.

** Catalyst 4040TM is a 40% solution of toluene sulfonic acid in isopropanol. In ink 29, catalyst 4040 was added to the activation solution.

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Ink Containing Resole Resin (no polymerized binder solution added):		
5	Ink Identification	30
10	Ingredient	Weight Fraction
	Polymerized Binder Solution S1	.000
	Hydroxyethyl Methacrylate	.2246
	Ethoxyethyl Methacrylate	.0254
	Methacrylic Acid	.0043
	Azobisisobutyronitrile	.0013
	Resole*	.4863
	Ethyl Lactate	.1553
	Phthalocyanine Blue	.0026
	Titanium Dioxide	.0531
	Catalyst 4040TM**	.0471
	<u>TOTAL</u>	<u>1.0000</u>

20 * Resole is a phenol-formaldehyde resin.

** Catalyst 4040TM is a 40% solution of toluene sulfonic acid in isopropanol added to activated ink paste.

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Preparation of Inks:

30 The ethyl lactate, phthalocyanine blue and titanium dioxide were ground together in a ceramic ball mill until the particle size was less than 5 microns on a Hegman grind gauge to form a ground pigment mixture. At the end of the grind cycle the polymerized binder solution (or resole if used) was added to the ground mixture to form a paste. The remaining ingredients termed the "activation solution" were mixed with the paste to form an "activated ink paste". The activation solution contains hydroxyethyl methacrylate, ethoxyethyl methacrylate, methacrylic acid, azobisisobutyronitrile, adhesion promoter i.e. CymelTM resin, and optionally an added catalyst such as Catalyst 4040TM. When a catalyst is employed it can be added to the activation solution or to the activated ink paste to form the finished ink termed "ink".

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EXAMPLES IN ACCORDANCE WITH THE FIRST ASPECT OF THE INVENTION

Lens Printing Procedure For Each Lens

45 The lenses were printed, i.e. a type of coating, with a colored dot pattern in accordance with U.S. Patent No. 4,582,402, (Knapp).

For each ink and lens the ink was placed on a metal cliche containing an image formed by a pattern of recessed cells. Excess ink was removed from the cliche by wiping the surface with a doctor blade. Ink in the recessed image was removed by transferring it to a soft rubber pad which was pressed over the 50 recessed image. Ink on the rubber pad was then transferred to a lens by contacting the inked part of the pad to a lens. Freshly printed lenses were heated up to 3 hours at 90°C in a nitrogen atmosphere.

Adhesion Testing Procedure

55 Printed lenses were hydrated for 2 hours at 98°C in a pH 8 saline solution. Hydrated lenses were examined to determine if any print pattern was removed during the hydration process. This is termed "Test 1". A more severe test, "Test 2", was performed by mounting the printed side of the hydrated lens four

inches from an air brush. The air brush was used to spray the lens for 30 seconds with methanol pressured at 40 pounds per square inch (psi) of nitrogen. After the air brush treatment, the lens was placed in pH 7.4 saline solution for 2-3 minutes.

The adhesion was considered not to pass (NP) if any coating came off during hydration in Test 1 or during the last saline treatment in Test 2. Otherwise the adhesion was considered to be passing (P). Adhesion test results are given below.

SUMMARY OF ADHESION TEST RESULTS FOR FIRST ASPECT OF THE INVENTION								
10	INK NO.	ADHESION PROMOTER		CATALYST LEVEL (%) [*]	ADHESION TEST P = PASS; NP = NOT PASS			
		Type	LEVEL IN INK		LENS POLYMER			
			(%)		L1		L2	
					Test 1	Test 2	Test 1	Test 2
15	1	CYMER	1.21	--	--	--	P	P
	2	301TM	9.39	--	--	--	P	P
	3		5.27	--	P	P	--	--
	4		2.54	0.42	P	P	--	--
20	5	CYMER	1.01	--	P	P	--	--
	6	327TM	9.07	--	P	P	--	--
	7		1.84	--	--	--	P	NP
	8		9.42	--	--	--	P	NP
	9		3.84	1.15	P	P	P	NP
	10		9.54	1.08	P	P	P	NP
	11		5.28	0.26	P	P	P	NP
	12		9.76	0.25	P	P	P	P
25	13	CYMER	1.19	0.27	P	P	P	NP
	14	385TM	9.08	0.25	P	P	P	NP
	15	CYMER	1.11	1.09	P	P	P	P
	16	1123TM	9.68	1.01	P	P	P	P
30	17	CYMER	1.20	0.99	--	--	--	--
	18	1141TM	9.22	1.33	--	--	--	--
	19	CYMER 1171TM	1.47	1.02	P	NP	P	NP
	20	CYMER	4.90	1.49	P	P	P	P
35	21	1172TM	9 ⁷⁷	2 ⁷⁸	I	I	I	I
	22		1.17	--	--	--	P	P
	23		4.91	--	--	--	P	NP
	24		9.58	--	--	--	P	NP
	25	BEETLE	2.35	--	--	--	P	P
40	26	60TM	5.04	--	--	--	P	NP
	27		9.07	--	--	--	P	NP
	28		3.09	--	P	NP	--	--
	29		3.97	0.42	P	P	--	--

* Catalyst 4040TM (40%) toluene sulfonic acid in isopropanol).

** The inks gelled before they could be printed.

55 The print pattern on all lenses passed Test 1. Most print patterns with lens polymer L1 passed Test 2 and a fewer number of lens polymers L2 passed Test 2. The testing procedure of Test 2 wherein the lenses are sprayed with methanol is physically and chemically very harsh -- more harsh than the normal rubbing to which the lenses would be subjected in normal use. Hence a lens which did not pass Test 2 does not

necessarily indicate a useless lens, since such lenses were found to at least pass Test 1.

The vast majority of the lenses containing the functional groups -OH and -COOH (i.e. those constructed of lens polymer L1) passed Test 2. Exceptions were a lens printed with ink containing Cymel 1171TM (ink no. 19), and another lens printed with ink containing Beetle 60TM (ink no. 28). It is believed that a higher amount of Cymel 1171TM or Beetle 60TM in the ink or a higher catalyst level would improve the result. At least one lens printed with inks containing the other additional compounds having the group -CH₂-O-R' passed the test.

As might be expected, as a group, more lenses that had functional groups i.e. L1 were found to pass both Test 1 and Test 2 than lenses that lack the functional groups -OH, -COOH, or -NH-R, i.e. lenses constructed of lens polymer L2. However at least one lens of the latter type passed the test when the additional compound in the ink was Cymel 301TM, Cymel 327TM, Cymel 1123TM, Cymel 1172TM, and Beetle 60TM. This indicates that the invention is indeed viable for hydrophilic lenses that do not have the functional groups -OH, -COOH, or -NH-R.

Both inks containing Cymel 1141TM gelled before they could be printed. It is believed that use of less catalyst, a different catalyst or no catalyst followed by overcoating the printed lens with a solution containing catalyst would overcome the gelling problem.

EXAMPLE IN ACCORDANCE WITH THE SECOND ASPECT OF THE INVENTION

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The lens printing procedure and adhesion testing procedure were similar as to those used in the Examples for the First Aspect of the Invention except that no polymerized binder solution S1 was added to the ink used for printing. The results are as follows:

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SUMMARY OF ADHESION TEST RESULTS FOR SECOND ASPECT OF THE INVENTION									
30	INK NO.	ADHESION PROMOTER		CATALYST LEVEL (%)	ADHESION TEST P = PASS; NP = NOT PASS				
		Type	LEVEL IN INK (%)		LENS POLYMER				
					L1		L2		
					TEST 1	TEST 2	TEST 1	TEST 2	
35	30	RESOLE	48.63	4.71	P	P	P	NP	

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The lens containing the functional groups -OH and -COOH (i.e., those constructed of lens polymer L1) passed both Test 1 and Test 2, while lens L2 without the functional groups -OH and -COOH only passed Test 1.

While the present invention has been described in conjunction with the specific embodiments set forth above, many alternatives, modifications and variations therefore will be apparent to those of ordinary skill in the art. All such alternatives, modifications and variations are intended to fall within the spirit and scope of the present invention.

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Claims

1. A method for making a colored contact lens comprising the steps of:
 - a) providing a contact lens constructed of non-hydrophilic polymers having functional groups selected from at least one of -COOH, -OH, and -NH-R, wherein R is hydrogen or C₁ to C₈ alkyl or hydrophilic polymers;
 - b) coating at least a portion of a surface of the lens with a color coat comprising at least one pigment, binding polymer having functional groups selected from at least one of -COOH, -OH, and -NH-R, wherein R is hydrogen or C₁ to C₈ alkyl, and an adhesion promotor having at least two functional groups per molecule of the formula -CH₂-O-R' wherein R' is hydrogen or C₁ to C₁₆ alkyl, and said -CH₂-O-R' groups are

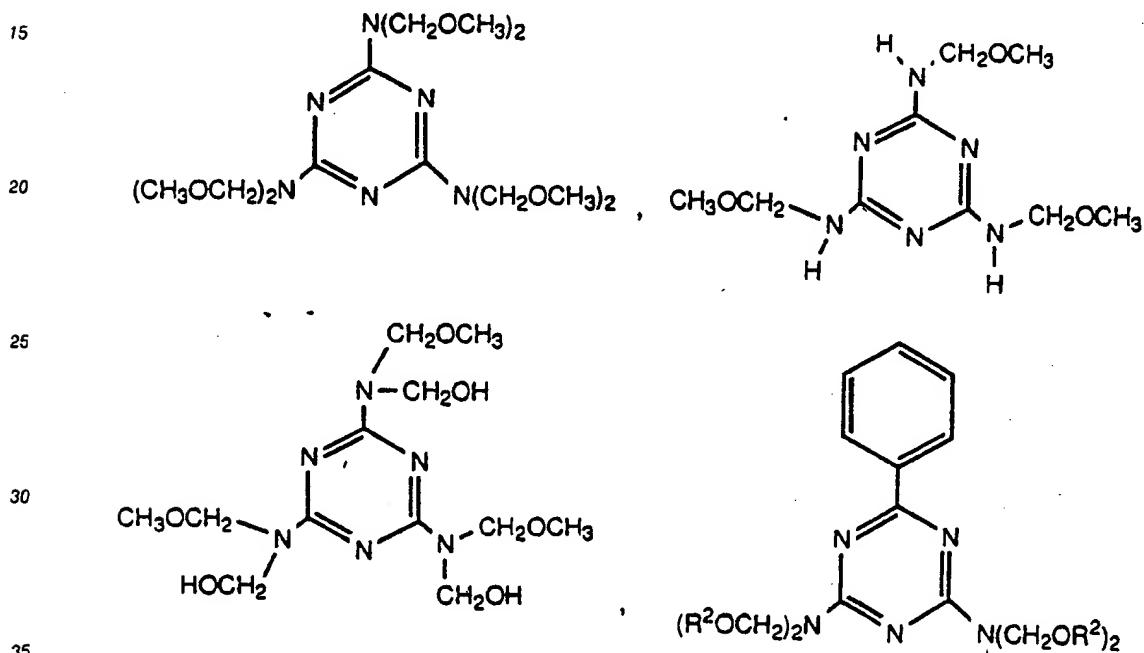
attached to a carbon atom that is part of an aromatic ring, or attached to a nitrogen or oxygen atom;
 c) subjecting the coated lens to conditions which cause the color coat to adhere to the lens.

2. The method of claim 1 wherein the lens polymer is hydrophilic and has functional groups selected from at least one of -COOH, -OH, and -NH-R, wherein R is hydrogen or C₁ to C₈ alkyl.

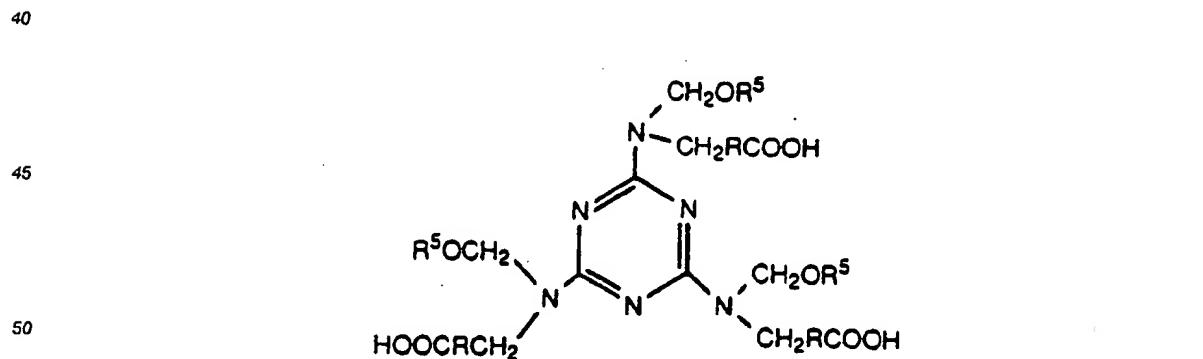
5 3. The method of claim 1 or claim 2 wherein the lens polymer and binding polymer are formed from a mixture of monomers comprising hydroxyethyl methacrylate, ethoxyethyl methacrylate, and methacrylic acid.

4. The method of claim 1 wherein the lens polymer is hydrophilic and is formed from monomers comprising vinyl pyrrolidone and hydrophobic monomer.

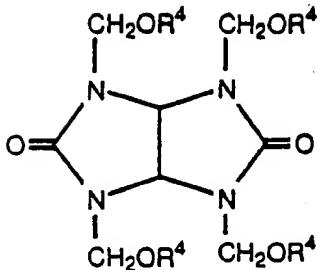
10 5. The method of any one of claims 1 to 4 wherein the adhesion promotor has the simplified structural formula



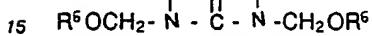
wherein R² is CH₃ or C₂H₅,



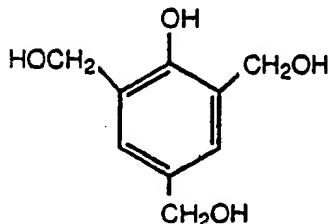
55 wherein R is a carbon-to-carbon single bond or C₁ to C₆ alkylene and R⁵ is independently C₁ to C₆ alkyl.



wherein each R⁴ is independently H, -CH₃, -C₂H₅, -C₃H₇, or -C₄H₉,



wherein each R⁶ is independently H, -CH₃, or isobutyl, or



polymers thereof and combinations thereof.

6. A lens made by the method of any one of claims 1 to 5.

30 7. A method for making a colored contact lens comprising the steps of:

- providing a contact lens constructed of non-hydrophilic polymers having functional groups selected from at least one of -COOH, -OH, and -NH-R, wherein R is hydrogen or C₁ to C₈ alkyl or hydrophilic polymers;
- coating at least a portion of a surface of the lens with a color coat comprising at least one pigment and adhesion promotor having functional groups of the formula -CH₂-O-R¹ wherein R¹ is hydrogen or C₁ to C₁₆ alkyl with said -CH₂-O-R¹ functional groups attached to a carbon atom that is part of an aromatic ring, or attached to a nitrogen or oxygen atom;
- subjecting the coated lens to conditions which cause the color coat to adhere to the lens.

35 8. A lens made by the method of claim 7.

40 9. A method for making a colored contact lens comprising the steps of:

- providing a contact lens constructed of polymer having functional groups of the formula -CH₂-O-R¹ wherein R¹ is hydrogen or C₁ to C₁₆ alkyl with said -CH₂-O-R¹ functional groups attached to a carbon atom that is part of an aromatic ring or attached to a nitrogen or oxygen atom;
- coating at least a portion of a surface of the lens with a color coat comprising at least one pigment and binding polymer having functional groups selected from at least one of the -COOH, -OH, and -NH-R, wherein R is hydrogen or C₁ to C₈ alkyl, and
- subjecting the coated lens to conditions which cause the color coat to adhere to the lens.

45 10. A lens made by the method of claim 9.

50 11. A method for making a colored contact lens comprising the steps of:

- providing a contact lens constructed of polymer having functional groups of the formula -CH₂-O-R¹ wherein R¹ is hydrogen or C₁ to C₆ alkyl with said -CH₂-O-R¹ functional groups attached to a carbon atom that is part of an aromatic ring or attached to a nitrogen or oxygen atom;
- coating at least a portion of a surface of the lens with a color coat comprising at least one pigment, binding polymer having functional groups of the formula -CH₂-O-R¹ wherein R¹ is hydrogen or C₁ to C₁₆ alkyl with said functional groups attached to a carbon atom that is part of an aromatic ring or attached to a nitrogen or oxygen atom, and an adhesion promotor having at least two groups per molecule selected from at least one of -COOH, -OH, and -NH-R, wherein R is hydrogen or C₁ to C₈ alkyl, and
- subjecting the coated lens to conditions which cause the color coat to adhere to the lens.

12. A lens made by the method of claim 11.
13. A colored contact lens comprising:
 - a) a lens constructed of polymer;
 - b) pigment mixed with adhesion promotor on at least a portion of a surface of the lens,
- 5 wherein the adhesion promotor and lens polymer are bonded to each other by the reaction of first functional groups selected from at least one of -COOH, -OH, and -NH-R, wherein R is hydrogen or C₁ to C₈ alkyl with second functional groups of the formula -CH₂-O-R' wherein R' is hydrogen or C₁ to C₁₆ alkyl with said second functional groups attached to a carbon atom that is part of an aromatic ring or attached to a nitrogen or oxygen atom.
- 10 14. The colored contact lens of claim 13 wherein the lens polymer is formed from at least one monomer selected from acrylic acid, methacrylic acid, hydroxy C₁ to C₈ alkyl ester of acrylic or methacrylic acid, amino C₁ to C₈ alkyl ester of acrylic or methacrylic acid, glycerol ester of acrylic or methacrylic acid, and mixtures thereof.
- 15 15. The lens of claim 13 wherein the lens polymer is formed from at least one monomer selected from acrylic acid, methacrylic acid, and hydroxyethyl methacrylate.

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(54) Color binding mechanism for contact lenses.

(57) Novel colored contact lens and methods for preparation thereof are described. Binding of a color coat to the lenses is accomplished by use of compounds containing the functional groups -CH₂-O-R¹, wherein R¹ is hydrogen or C₁ to C₁₆ alkyl, and said -CH₂-O-R¹ groups are attached to a carbon atom that is part of an aromatic ring, or are attached to a nitrogen or oxygen atom.

EP 0 393 532 A3



DOCUMENTS CONSIDERED TO BE RELEVANT			CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	
A,D	EP-A-0 187 137 (SCHERING) * Claim 1 * ---	1	C 08 J 7/04 G 02 B 1/10
A	US-A-4 490 495 (H. WEBER) * Claim 1; column 4, lines 18-61 * ---	1	
A	US-A-4 127 697 (B.L. LAURIN) * Claim 1 * ---	1	
A	US-A-4 127 682 (B.L. LAURIN) * Claim 1 * ---	1	
D,A P	US-A-4 857 072 (K.W. NARDUCY) * Claim 1 * -----	1	
			TECHNICAL FIELDS SEARCHED (Int. Cl.5)
			C 08 J G 02 B
<p>The present search report has been drawn up for all claims</p>			
Place of search	Date of completion of the search		Examiner
THE HAGUE	18-09-1990		DEPIJPER R.D.C.
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